

Claim Amendments

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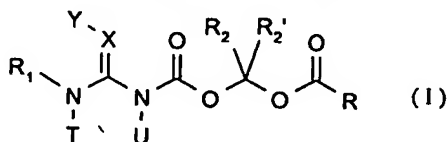
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4/20/07

Claims 1-27 (Cancelled)

Claim 28. (New) A compound of formula (I)



wherein

R<sub>1</sub> thiazolyl-substituted C<sub>1</sub>-C<sub>6</sub>alkyl, wherein the thiazolyl moiety is unsubstituted or mono- or poly-substituted by identical or different halogen atoms;

X is CH;

Y is NO<sub>2</sub>;

T is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl;

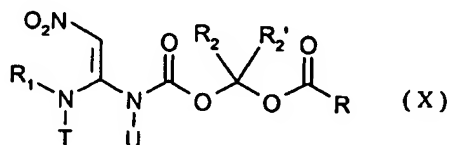
U is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl;

R<sub>2</sub> is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl;

R<sub>2</sub>' is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl; and

R is C<sub>1</sub>-C<sub>20</sub>alkyl, being unsubstituted or substituted by one or more identical or different substituents, the said substituents being selected from the group halogen, cyano, nitro, hydroxy, C<sub>1</sub>-C<sub>6</sub>alkoxy, C<sub>1</sub>-C<sub>6</sub>alkylthio, C<sub>1</sub>-C<sub>6</sub>haloalkyl, C<sub>1</sub>-C<sub>6</sub>haloalkoxy and phenyl.

Claim 29. (New) A compound according to claim 28, which is a compound of formula (X)



wherein

R<sub>1</sub> is --CH<sub>2</sub>-thiazolyl, which is unsubstituted or mono- or poly-substituted by identical or different halogen atoms;

R is C<sub>1</sub>-C<sub>20</sub>alkyl, being unsubstituted or mono- or poly-substituted by identical or different substituents, the said substituents being selected from the group halogen, cyano, nitro, hydroxy, C<sub>1</sub>-C<sub>6</sub>alkoxy, C<sub>1</sub>-C<sub>6</sub>alkylthio, C<sub>1</sub>-C<sub>6</sub>haloalkyl, C<sub>1</sub>-C<sub>6</sub>haloalkoxy and phenyl;

T and U are each independently of the other hydrogen, methyl or ethyl;

R<sub>2</sub> is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl; and

R<sub>2</sub>' is hydrogen or C<sub>1</sub>-C<sub>6</sub>alkyl.

Claim 30. (New) A compound of formula (I) according to claim 28, wherein U is methyl or ethyl.

Claim 31. (New) A compound of formula (I) according to claim 28, wherein T is methyl or ethyl.

Claim 32. (New) A compound of formula (I) according to claim 28, wherein R<sub>2</sub>, R<sub>2</sub>' are each independently of the other hydrogen, methyl, or ethyl.

Claim 33. (New) A compound of formula (I) according to claim 28, wherein R<sub>1</sub> is —CH<sub>2</sub>—thiazolyl, that is unsubstituted or mono- or di-substituted by halogen.

Claim 34. (New) A compound of formula (I) according to claim 33, wherein thiazolyl in R<sub>1</sub> is 2-chlorothiazol-5-yl.

Claim 35. (New) A compound according to claim 28, wherein R is straight-chain or branched C<sub>6</sub>-C<sub>20</sub>alkyl.

Claim 36. (New) A preparation for controlling parasites on warm-blooded animals, comprising a compound according to claim 28, one further parasiticide, and a physiologically tolerable carrier.

Claim 37. (New) A parasiticial composition comprising a compound of formula (I) according to claim 28 and at least one physiologically tolerable carrier.

Claim 38. (New) A parasiticial composition according to claim 37, comprising from 0.1 to 99% by weight of a compound of formula (I) according to claim 1 and from 99.9 to 1% by weight of a solid or liquid, physiologically tolerable carrier, including from 0 to 25% by weight of a non-toxic dispersant.

Claim 39. (New) A parasiticial composition according to claim 38, which is a pour-on or spot-on formulation.

Claim 40. (New) A method of controlling parasites on warm-blooded animals, which comprises administering to a warm blooded animal a parasitically effective compound of formula (I) according to claim 28.

Claim 41. (New) A method according to claim 40 comprising the topical application of a compound of formula (I) according to claim 28.

Claim 42. (New) A method according to claim 40, wherein a compound of formula (I) according to claim 1 is administered in a dose of from 0.01 to 800 mg/kg/body weight.

Claim 43. (New) A veterinary medicinal preparation against parasites comprising a compound of claim 28<sub>u</sub>

and a physiologically tolerable carrier.